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ORIGINAL ARTICLE

Pelvic muscle floor rehabilitation as a therapeutic option in lifelong premature ejaculation: long-term outcomes

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The aim of the study was to evaluate the long-term outcomes of pelvic floor muscle (PFM) rehabilitation in males with lifelong premature ejaculation (PE), using intravaginal ejaculatory latency time (IELT) and the self-report Premature Ejaculation Diagnostic Tool (PEDT) as primary outcomes. A total of 154 participants were retrospectively reviewed in this study, with 122 completing the training protocol. At baseline, all participants had an IELT ≤ 60 s and PEDT score >11. Participants completed a 12-week program of PFM rehabilitation, including physio-kinesiotherapy treatment, electrostimulation, and biofeedback, with three sessions per week, with 20 min for each component completed at each session. The effectiveness of intervention was evaluated by comparing the change in the geometric mean of IELT and PEDT values, from baseline, at 3, 6, and 12 months during the intervention, and at 24 and 36 months postintervention, using a paired sample 2-tailed *t*-test, including the associated 95% confidence intervals. Of the 122 participants who completed PFM rehabilitation, 111 gained control of their ejaculation reflex, with a mean IELT of 161.6 s and PEDT score of 2.3 at the 12-week endpoint of the intervention, representing an increase from baseline of 40.4 s and 17.0 scores, respectively, for IELT and PEDT (P < 0.0001). Of the 95 participants who completed the 36-month follow-up, 64% and 56% maintained satisfactory ejaculation control at 24 and 36 months postintervention, respectively. *Asian Journal of Andrology* (2018) **20**, 572–575; doi: 10.4103/aja.aja_30_18; published online: 3 July 2018

Keywords: biofeedback; electrostimulation; intravaginal ejaculatory latency time; pelvic floor rehabilitation; premature ejaculation; premature ejaculation diagnostic tool

INTRODUCTION

Premature ejaculation (PE) is a common sexual dysfunction in males which has a significant impact on the quality of life of the patients and partners.^{1,2} The role of the pelvic floor muscles in controlling the ejaculation is often overlooked in patients with PE.³

PE can either be lifelong or acquired. The 2014 updated criteria of the International Society for Sexual Medicine (ISSM) defines lifelong PE as ejaculation usually occurring within 1 min of vaginal penetration, and acquired PE as a clinically significant reduction in intravaginal ejaculatory latency time (IELT), usually in the range ≤ 3 min or less, inability to delay ejaculation and negative personal consequences, such as stress, bother, frustration, and/or the avoidance of sexual intimacy.^{4,5} Among factors contributing to acquired PE, IELT is defined as the time from vaginal intromission to intravaginal ejaculation⁴ and is often used as a parameter to quantify clinical response to therapy and as a standardized method to compare different treatment modalities in clinical trials.

In general, patients with PE are not aware of the role of pelvic floor muscles in the control of ejaculation.^{3,6} Therefore, it is possible that in many of these patients, PE may not result from an early arrival of the

stimulus but, rather, due to a failure to selectively produce an effective contraction of the pelvic floor.⁷ Therefore, in our study, men with lifelong PE underwent pelvic floor muscle (PFM) rehabilitation, using a modification of techniques used for the treatment of urinary and fecal incontinence,⁶ and including physio-kinesiotherapy treatments, electrostimulation, and biofeedback.

The primary endpoint of our study was to evaluate the effectiveness of PFM rehabilitation by measuring the change, from baseline, in IELT and the self-administered Premature Ejaculation Diagnostic Tool (PEDT) over a follow-up period of 36 months. The secondary endpoint was to investigate the long-term effectiveness of PFM rehabilitation in training patients to recognize when and how to control the muscles involved in the ejaculatory reflex.

PARTICIPANTS AND METHODS

Participants

A total of 154 male participants were retrospectively reviewed in this study, which was conducted between September 2012 and December 2013. To be included in the study, participants had to be in a stable relationship, with the same partner, for at least 6 months and to engage

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in sexual intercourse ≥ 1 per week. All participants provided written informed consent. Among the 154 participants enrolled into the study, 122 (79.2%) completed the training protocol, with the other 32 (20.8%) being excluded due to signs of prostatitis or personal preference to manage the problem with medication.

The study was conducted in accordance with the Ethical Principles for Medical Research Involving Human Subjects (World Medical Association, The Declaration of Helsinki Principles, 2000) and was approved by the local medical ethical committee of the hospital (ASL Lt/No. 43001/9674/2012).

PE was diagnosed by applying the ISSM definition of lifelong PE,⁵ with all participants having a baseline IELT of ≤ 60 s and PEDT test score >11. During the first visit, participants and their partners were interviewed individually and each was requested to provide an independent estimation of IELT. Participants also completed the PEDT, a self-administered questionnaire developed to standardize the diagnosis of PE and to capture the main elements of the DSM-IV-TR not considered by IELT (such as control, frequency, minimal sexual stimulation, distress, and interpersonal difficulty).⁸⁻¹⁰

All participants had previously tried different types of therapy, including esthetic creams, serotonergic antidepressants, and phosphodiesterase type 5 inhibitors (PDE5i), without a satisfactory response, especially with regard to adverse effects. Specifically, 47 participants had previously applied a local esthetic cream (EMLA 2.5 g, AstraZeneca, Cambridge, UK) to the glans of the penis and penile shaft, 30 min before sexual intercourse, with no perceived significant benefit with regard to IELT, and with numbness of penis reported as the most frequent adverse event. Nineteen participants had been treated using tramadol hydrochloride, a narcotic-like pain reliever. Twenty-seven participants had been treated using selective serotonin reuptake inhibitor (SSRI) as needed, with nausea and cost of the drug being the main causes of treatment dropout. Another 19 participants were treated with PDE5i as needed (sildenafil 50 mg or vardenafil 20 mg), 1 h to 3 h before sexual intercourse. All participants discontinued previous therapies at least 6 months before starting the program of PFM rehabilitation.

Preintervention IELT and PEDT were measured over a 4-week baseline period, with participants provided with a stopwatch and instructions on how to measure IELT and how to complete the PEDT. During this 4-week baseline period, participants were asked to have coitus at least four times. Couples were instructed not to use condoms or any topical esthetic cream and not to pause during intercourse or have interrupted intromission. Furthermore, participants were instructed to measure IELT only for the first intercourse, if intercourse took place more than once in a single session.

All participants underwent a physical examination prior to the start of the intervention. None of the participants had phimosis (19 patients were circumcised), frenulum breve, history of chronic prostatitis, or erectile dysfunction. The mean International Index of Erectile Function (IIEF) score of the study group prior to the intervention was 28.5 (95% confidence interval [CI]: 26.2–28.7; standard deviation [s.d.] = 1.24). Participants also underwent a urological screening, which included the Meares–Stamey test, to exclude the presence of bacterial prostatitis, and a digital rectal examination.¹¹

All participants completed the same PFM rehabilitation intervention,^{12,13} which consisted of the following components: physiokinesiotherapy to achieve a muscle contraction which provided participants with awareness of the motor activity; electrostimulation to the pudendal nerve for contraction of the puborectalis muscle, which causes the urethral sphincter to contract; and biofeedback to teach participants to control the muscle contractions of the perineal floor and the genitourinary sphincter.^{14,15} During the physiokinesiotherapy portion of the session, patients performed personalized physical exercises focused on isometric and isotonic contractions of the PFMs. Electrostimulation of the pudendal nerve, performed to enhance strengthening of the PFMs, was applied using a cylindrical anal probe, with contact between the electrodes in the anal probe and the anterior portion of the anal sphincter to stimulate the pubovisceral muscles (puborectal and pubourethral), with mild, painless electrical pulses used for stimulation.^{14,15}

Participants completed three 60-min sessions per week, during which the three techniques were applied for a period of 20-min each. Outcome measures were obtained after the first 18 sessions (6 weeks) and then again at the end of the intervention (12 weeks).

Effectiveness of PFM rehabilitation was evaluated through comparison of the group mean IELT values and PEDT scores at the 3-, 6-, and 12-month time point of treatment, and at the 24- and 36-month time point of follow-up. A successful outcome of the intervention was defined as ability to control the ejaculatory reflex and a PEDT score ≤ 8 .^{9,10}

Statistical analyses

The paired sample 2-tailed *t*-test, with the associated 95% CI, was used to compare the geometric mean of IELTs and PEDT at each time point of measurement, with P < 0.05 being considered to be statistically significant. Statistical analyses were conducted using SPSS (version 20.0, SPSS Inc., Chicago, IL, USA) and JMP (version 10.0, SAS Institute Cary, NC, USA).

RESULTS

The 122 participants included in the final analysis had a mean age of 24 years (range, 18–41 years), and a mean baseline IELT and PEDT of 40.4 s (95% CI: 39.9–40.9) and 17.0 (95% CI: 16.7–17.3), respectively. At the end of the 12-week PFM rehabilitation program, 111 (90.9%) of the 122 patients reported a satisfactory ejaculation latency time from the start of intravaginal intercourse, with a mean IELT and PEDT of 161.6 s (95% CI: 159.0–164.2) and 2.3 (95% CI: 2.1–2.5), respectively (**Table 1**).

All participants were followed up for at least 2 years after the intervention, with 95 of the 122 participants (77.8%) completing the follow-up to 36-month postintervention. The mean follow-up period was 36.4 months (range, 28.2–43.6 months), and satisfactory ejaculation latency time were maintained throughout the follow-up period (at least 2 years following the 12-week intervention) in 64% of participants. Both

Table 1: Mean intravaginal ejaculatory latency time and premature ejaculation diagnostic tool scores and rate of satisfied patients after pelvic floor muscle rehabilitation. The paired sample two-tailed *t*-test with associated 95% confidence intervals was used to compare the geometric mean of intravaginal ejaculatory latency time and premature ejaculation diagnostic tool

	Mean IELT (s.d.)	Mean PEDT (s.d.)	Patient evaluated (n)	Satisfied patient (%)
Baseline	40.4 (8.7)	17.0 (2.2)	122	
At 3 months	161.6 (14.5)	2.3 (1.1)	122	90.2
At 6 months	157.7 (12.9)	2.8 (1.3)	122	85.3
At 12 months	156.3 (13.1)	3.4 (1.6)	122	78.5
At 24 months	149.0 (12.2)	4.6 (1.0)	122	63.9
At 36 months	138.5 (10.5)	5.7 (1.2)	95	56.8
Р	< 0.0001	< 0.0001		

IELT: intravaginal ejaculatory latency time; PEDT: premature ejaculation diagnostic tool; s.d.: standard deviation



IELT and PEDT were improved at the endpoint of follow-up, compared to baseline (IELT, P < 0.0001; **Table 1** and **Figure 1**; PEDT, P < 0.0001; **Table 1** and **Figure 2**). Compared to baseline, IELT improved by 40.4 s and the PEDT by a score of 17.04 (P < 0.0001).

Of note, at the 3-month time point of measurement, 11 (9.0%) participants did not report any significant improvement in ejaculatory latency time. At the 2-year time point of follow-up, 27 (22.1%) of the 122 patients who reported a significant improvement in clinical condition after PFM rehabilitation voluntarily dropped out of the study. At 24- and 36-month time point of follow-up, 63.9% and 56.8% of participants, respectively, maintained satisfactory ejaculation control (**Table 1**). No adverse effect was reported that participants could have led to the discontinuation of the PFM rehabilitation.

DISCUSSION

Currently, the recommended management of PE includes behavioral and psychotherapy, drug therapy, and/or a combination of these.¹⁶⁻¹⁹ Pharmacotherapy, such as dapoxetine (a short-acting SSRI that is the only approved pharmacological treatment for premature ejaculation), used as necessary, or other off-label antidepressants (daily SSRIs), which require regular dosage, have the highest level of evidence of effectiveness.¹⁶⁻¹⁹ Various other forms of pharmacotherapy have been used for the treatment of PE, including topical local anesthetics, tramadol, PDE5 inhibitors, and α -adrenergic blockers, each being supported by varying levels of clinical evidence.¹⁶⁻¹⁹ Research evidence has supported a possible active role of the PFMs, particularly the ischiocavernosus and bulbocavernosus muscles and sphincters, in the control of ejaculation, as confirmed by a significant increase in electromyography activity during the entire physiological process.²⁰ As we have previously reported,^{15,21} physiokinesiotherapy and electrostimulation are designed to improve the contractile strength of the perineal muscles, whereas biofeedback is used to facilitate patients' learning to recognize and contract PFMs to increase the closing strength of the urethral sphincter. However, PFM training does require a few months for a patient to understand the dynamics of the sequence of events, learn to control the ejaculatory reflex and naturally apply this learning during sexual intercourse.^{11,21} In general, twenty sessions of training are required for a patient to gain sufficient control over the ejaculatory reflex. It is reasonable to assume that some patients may require a faster therapeutic approach, such as drug therapy. Another limitation of training is that not all patients are able to perform a selective control of the pelvic floor muscles or recognize the sensation that precedes the inevitability of ejaculatory reflex. However, when PFM rehabilitation is performed adequately, patients could benefit from PE control for 6 to 36 months, without the use of drugs.²¹ The findings of the current study confirm our previous hypothesis regarding the clinical benefit of PFM rehabilitation in allowing patients to achieve pelvic floor awareness and improved self-confidence and sense of control over their ejaculatory reflex.²¹ In the current study,

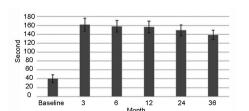


Figure 1: IELTs evaluated at 3, 6, 12, 24, and 36 months after PFM rehabilitation. IELT: intravaginal ejaculatory latency time; PFM: pelvic floor muscle.

the improvement in IELT, from baseline, of 146.2 ± 38.3 s was greater after 12 weeks of training than in our previous study.²¹ Furthermore, compared to baseline, IELT and PEDT were found to be significantly improved among those patients who completed the follow-up to 24 (63.9%) and 36 (56.8%) months postintervention. These results represent another important achievement obtained by way of an easily learned technique that can be mastered using pelvic floor biofeedback. Moreover, no adverse effects of the PFM rehabilitation protocol were identified, compared to other medical therapies such as gastrointestinal symptoms (nausea and diarrhea) and dizziness and headaches, which have been associated with the use of dapoxetine.^{22,23} In addition, the long-term effects of dapoxetine on reproductive functions should be cleared as some recent data demonstrated its negative impact on fertility.²⁴

To the best of our knowledge, the present study has the longest follow-up period (36 months) reported to date for any intervention for PE. However, several limitations of our study should be noted. The main limitation regards the lack of a control group. It is important to note that we planned our study without a control arm as we consider that it is unethical to not treat patients with PE when it impacts quality of life. We also used the PEDT test as a primary outcome measure, although the Italian version of the PEDT remains to be validated. Furthermore, we used the PEDT as a measure of the impact of PFM training on PE, noting that the PEDT was designed to screen for PE rather than as patient-reported outcome for an intervention. In addition, we did not consider the perception of orgasmic intensity in our study, which has previously been reported in men with PE.25 Future studies should consider using the validated "Orgasmometer," which is a psychometric tool that evaluates the subjective perception of orgasmic intensity²⁵ on a 10-point scale. There is a lack of standardized measures to assess improvement following the PFM intervention, with the treatment parameters difficult to define as they are largely based on sensory perception. Nonetheless, there is a continued need for evidence-based research to validate the role of physiokinesiotherapy in the treatment of PE. Although the type, the amount, and the focus (relaxation, strength, support, or control) of exercise required have not yet been standardized, the long-term results obtained in our patients with lifelong PE suggest that it may be considered as a valid therapeutic option for patients with PE.

CONCLUSION

The pelvic floor muscle rehabilitation protocol is easy to perform, with no reported adverse effects. Although it has not yet been standardized, the long-term results obtained in our patients with lifelong premature ejaculation suggest that it may be considered as a therapeutic option for patients with premature ejaculation. However, controlled trials are needed before making final evaluation on the effectiveness of pelvic floor muscle rehabilitation in premature ejaculation patients.

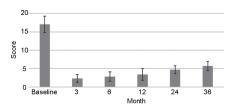


Figure 2: PEDT scores evaluated at 3, 6, 12, 24, and 36 months after PFM rehabilitation. PEDT: premature ejaculation diagnostic tool; PFM: pelvic floor muscle.

AUTHOR CONTRIBUTIONS

ALP, GP, YAS, AF, GB, EI, AZ: Study conception and design, data collection and statistical analyses; ALP, GP, YAS, AF, AC; EC: Writing and revision of the manuscript.

COMPETING INTERESTS

All authors declared no competing interests.

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